## P/ TT COOPERATION TREAT'

	From the INTERNATIONAL BUREAU	
PCT~	To:	
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NOTIFICATION OF ELECTION	United States Patent and Trademark	
(PCT Rule 61.2)	(Box PCT) Crystal Plaza 2 Washington, DC 20231 États-Unis d'Amérique	
Date of mailing (day/month/year) 07 October 1998 (07.10.98)	in its capacity as elected Office	
International application No. PCT/US98/02899	Applicant's or agent's file reference 2833.58	
International filing date (day/month/year) 13 February 1998 (13.02.98)	Priority date (day/month/year) 13 February 1997 (13.02.97)	
Applicant	and the second s	
MAZESS, Richard, B. et al		
The designated Office is hereby notified of its election made.	de:	
X in the demand filed with the International Preliminal	ry Examining Authority on:	
14 Septembe	r 1998 (14.09.98)	
·	<del></del>	
in a notice effecting later election filed with the Inter	national Bureau on:	
2. The election X was		
was not		
made before the expiration of 19 months from the priority Rule 32.2(b).	date or, where Rule 32 applies, within the time limit under	
•		
·	•	
	·	
	Authorized officer	
The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland	Authorized officer  Nicola Wolff	
Facsimile No.: (41-22) 740.14.35	Telephone No.: (41-22) 338.83.38	

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connector, via covalent bonding, hydrogen bonding, metallic bonding, van der Wall forces, ionic bonding, coulombic forces, hydrophobic or hydrophilic forces, adsorption or absorption, chelate type association, or any combination thereof. The terms "moiety" and "component" used in connection with vitamin D or D, or in connection with target molecule moiety or T, are meant to refer to vitamin D or target molecule in the conjugated forms disclosed herein, i.e., after association occurs. Association between the vitamin D analog and the target molecule may occur at any position on the vitamin D analog molecule depending on the functionality of the target molecule. For example, a bisphosphonate or amide may suitably link at positions on the vitamin D compound or vitamin D analog molecule having a hydroxyl group, such as at C-1, C-3, C-24, C-25.

Vitamin D compounds and analogs operable in the present invention are suitably represented by formula (II):

$$\begin{array}{c|c} & & \\ \hline \\ & & \\ \\ & & \\ \\ & & \\ \\ & & \\ \end{array}$$

wherein  $R^1$  is H or OH; Z represents a saturated or unsaturated, substituted or unsubstituted, straight-chain or branched  $C_1$  -  $C_{18}$  hydrocarbon group; Y is a =  $CH_2$  group; and t is 0 or 1, such that when t is 0, the compound of formula (II) is a 19-nor compound. Preferably, Z is a side chain represented by formula (IIIA):

$$R^{3}$$
  $R^{4}$   $X$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$ 

wherein  $R^1$  is H or OH;  $R^2$  is H or OH;  $R^3$  is  $CH_3$  or H;  $R^4$  is H or OH, X is O or S; Y is NH, O or NR wherein R is H or  $C_1$ - $C_4$  alkyl; n is an integer from 1 to 4;  $R^8$  and  $R^9$  are each H or taken together form a double bond between C-22 and C-23; and pharmaceutically acceptable salts thereof, i.e., the bisphosphonate is linked at the C-25 position of the vitamin D moiety.

Also provided are conjugates of formula (X):

wherein  $R^1$  is H or OH;  $R^2$  is H or OH;  $R^3$  is  $CH_3$  or H;  $R^4$  s H or OH, X is O or S; Y is NH, O or NR wherein R is H or  $C_1$ - $C_4$  alkyl;  $R^5$  is H or OH; n is an

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integer from 1 to 4; R<sup>8</sup> and R<sup>9</sup> are each H or taken together form a double bond between C-22 and C-23; and pharmaceutically acceptable salts thereof, i.e., the bisphosphonate is linked to the vitamin D moiety at C-3.

-Also provided are conjugates of formula (XI):

$$R^{3}$$
  $R^{4}$   $R^{5}$   $R^{5}$   $R^{6}$   $R^{7}$   $R^{8}$   $R^{8}$   $R^{8}$   $R^{8}$   $R^{8}$   $R^{8}$   $R^{8}$   $R^{9}$   $R^{8}$   $R^{5}$   $R^{5}$   $R^{5}$   $R^{5}$   $R^{6}$   $R^{7}$   $R^{7$ 

wherein  $R^2$  is H or OH;  $R^3$  is  $CH_3$  or H;  $R^4$  is H or OH, X is O or S; Y is NH, O or NR wherein R is H or  $C_1$ - $C_4$  alkyl;  $R^5$  is H or OH, n is an integer from 1 to 4,  $R^8$  and  $R^9$  are each H or taken together form a double bond between C-22 and C-23; and pharmaceutically acceptable salts thereof, i.e., the bisphosphonate linkage is at C-1 of the vitamin D moiety.

It is noted that typically the linkage between the bisphosphonate moiety and the vitamin D moiety is through a hydroxyl on the vitamin D where the hydroxyl is converted to a

group and is linked to the amine or hydroxy group, i.e., Y, of the bisphosphonate to form a carbamate-type or carbonate-type linkage. X can be O or S. For example, a hydroxyl group may be contained in the vitamin D structure at C-1, C-3, C-24, C-25, and conjugation can be effected at any hydroxyl position but is suitably one of the above.

09/402636-5000

### PATENT COOPERATION TREATY

## **PCT**

REC'D	15	OCT	1999
WIP	5	F	PCT

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

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Applicant's or agent's file reference		See Notification of Transmittal of International					
MFF/FP5726708 FOR FURTHER ACTION Preliminary Examination Report (Form PCT/IPEA/416)							
International application No.	International filing date (day/month/year) Priority date (day/month/year)						
PCT/US98/02899	13/02/1998	13/02/1997					
International Patent Classification (IPC) or I	national classification and IPC						
		,					
Applicant							
BONE CARE INTERNATIONAL, II	NC. et al.						
This international preliminary exa	mination report has been prepared to	by this International Preliminary Examining Authority					
and is transmitted to the applican	according to Article 36.						
2. This REPORT consists of a total	of 6 sheets, including this cover she	eet.					
been amended and are the b	ied by ANNEXES, i.e. sheets of the asis for this report and/or sheets co 607 of the Administrative Instruction	description, claims and/or drawings which hav ntaining rectifications made before this Authority ns under the PCT).					
These annexes consist of a total	of 3 sheets.						
3. This report contains indications re	elating to the following items:						
I ⊠ Basis of the report							
II □ Priority							
III   Non-establishment o	f opinion with regard to novelty, inve	entive step and industrial applicability					
IV ☐ Lack of unity of inver	ntion						
V   Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement							
VI ⊠ Certain documents o	cited						
VII Certain defects in the	international application						
VIII 🛛 Certain observations	on the international application	·					
L							

Date of submission of the demand

14/09/1998

Name and mailing address of the international preliminary examining authority:

European Patent Office
D-80298 Munich
Tel. +49 89 2399 - 0 Tx: 523656 epmu d
Fax: +49 89 2399 - 4465

Date of completion of this report

Authorized officer

Stoltner, A

Telephone No. +49 89 2399 8408

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US98/02899

I. Basis	of the	report
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1. This report has been drawn on the basis of (substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.):

	Des	cription, pages:					
	1-10 18-3	),12-15, 35	as originally filed				
	11,1	16,17	as received on	07/01/1999	with letter of	04/01/1999	
	Clai	ims, No.:					
	1-40	)	as originally filed				
	Dra	wings, sheets:					
	1/9-	9/9	as originally filed				
				•	•	·	
2.	The	amendments hav	e resulted in the cancellation	n of:			
		the description,	pages:				
		the claims,	Nos.:			•.	
		the drawings,	sheets:				
3.		This report has b considered to go	een established as if (some beyond the disclosure as file	of) the amendmer ed (Rule 70.2(c)):	nts had not been n	nade, since they have bee	η
4.	Add	ditional observation	ns, if necessary:			•	

V. R asoned stat m nt und r Articl 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: C

Claims

No:

Claims 1-40

Inventive step (IS)

Yes: Claims

No: Claims

Industrial applicability (IA)

Yes: Claims

1-40 (except claims 23-27, 37 and 39 for some contracting states

within the EPO)

No: Claims

2. Citations and explanations

see separate sheet

#### VI. Certain documents cited

1. Certain published documents (Rule 70.10)

and / or

2. Non-written disclosures (Rule 70.9)

see separate sheet

### VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

#### ad section V:

- 1). The present application concerns a conjugate for the targeted therapeutic delivery of vitamin D compounds to specific tissue, said conjugate comprising a vitamin D compound bound to a target molecule moiety as set out in the present claim 1.
- 2). The following documents are considered relevant for the subject-matter of the present application:
- D1, US-A-5 691 328, an intermediate document, discloses conjugates of vitamin D compounds with phosphoethanolamine with anti-tumor activity (cf. abstract) or for the treatment of osteomalacia (cf. col. 6, lines 37-47). As vitamin D compounds are bone therapeuting agents, it has also to be submitted that they are in some way "bone seeking" molecules, acting with affinity at a "tissue of interest" as pointed out in the application on page 8, last para. bridging with page 9, lines 1-7.
- D2, US-A-5 232 836, refers to vitamin D derivatives covalently bound to immunogenic carrier proteins and antibodies (cf. abstract). In D2, the problem of delivering vitamin D conjugates to specific target tissues is also addressed (cf. col. 2, last para., col. 3, lines 58-64). Moreover, D2 also discloses connecting groups by which to couple the vitamin D compound to the carrier (cf. col. 27, lines 10-35). As set out in the application on page 9, lines 27-30, the vitamin D derivatives coupled to antibodies have the effect of targeting the vitamin D to, e.g., tumors. Moreover, in his letter of 12/4/99, the Applicant admits that vitamin D indeed has "traditional target tissues".
- D3, WO-A-9 307 883, provides nucleosides linked to vitamin D by an aminoalkoxy or aminoalkylamino moiety (cf. claims 1-5). D3 encompasses the application of antisense agents for diagnostics and therapeutics where said antisense oligonucleotides are to be transported across cell membranes or "taken up by cells to express" activity. According to the Applicant, said cells have to be construed as pertaining to a "tissue of interest" (cells of interest).

## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

- D4, J. Pharmacol. Exp. Ther., 237(3), 1986, pp. 837-840, J.M. Landowski et al., publishes the conjugation of-vitamin D3 compounds to glycopyranosides and the administration of said conjugates to vitamin D deficient rats maintained on a low calcium diet, whereby an increase of the intestinal calcium transport as well as the mobilization of the bone calcium was observed. As in the present application the term "target molecule" also encompasses molecules that "influences metabolism of the tissue of interest" (cf. page 9, lines 12-13), the raise of intestinal calcium transport implicitly also covers the metabolism of a tissue of interest (e.g. the metabolism of fatty acids).
- 3). In view of the above, it has to be noted that a conjugate or a composition as specified in claims 1 and 20 is known in the prior art. As appending to the main claims 1 and 20, the subject-matter of the subsequent claims 2-19 and 21-40 has also to be considered as anticipated by the prior art teaching.
- 4). For the assessment of the present claims 23-27, 37 and 39 on the question whether they are industrially applicable, no unified criteria exist in the PCT. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.

### ad section VI:

1). US-A-5 691 328, with a priority date of 27/8/96 and a publication date of 25/11/97;

### ad section VIII:

1). The expression "..having an affinity for a tissue of interest" in claim 1 is unclear and includes no clear delimitation from the prior art. Moreover, this term can be construed as an attempt to define the invention by the result to be achieved, which gives no contribution to novelty or inventive step.

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US98/02899

2). Incorporations made by reference (cf. page 20, line 16) are not accepted in the working practice of the EPO.

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connector, via covalent bonding, hydrogen bonding, metallic bonding, van der Wall forces, ionic bonding, coulombic forces, hydrophobic or hydrophilic forces, adsorption or absorption, chelate type association, or any combination thereof. The terms "moiety" and "component" used in connection with vitamin D or D, or in connection with target molecule moiety or T, are meant to refer to vitamin D or target molecule in the conjugated forms disclosed herein, i.e., after association occurs. Association between the vitamin D analog and the target molecule may occur at any position on the vitamin D analog molecule depending on the functionality of the target molecule. For example, a bisphosphonate or amide may suitably link at positions on the vitamin D compound or vitamin D analog molecule having a hydroxyl group, such as at C-1, C-3, C-24, C-25.

Vitamin D compounds and analogs operable in the present invention are suitably represented by formula (II):

wherein  $R^1$  is H or OH; Z represents a saturated or unsaturated, substituted or unsubstituted, straight-chain or branched  $C_1$  -  $C_{18}$  hydrocarbon group; Y is a =  $CH_2$  group; and t is 0 or 1, such that when t is 0, the compound of formula (II) is a 19-nor compound. Preferably, Z is a side chain represented by formula (IIIA):

$$R^{3}$$
  $R^{4}$   $X$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$   $PO_{3}H_{2}$ 

wherein  $R^1$  is H or OH;  $R^2$  is H or OH;  $R^3$  is  $CH_3$  or H;  $R^4$  is H or OH, X is O or S; Y is NH, O or NR wherein R is H or  $C_1$ - $C_4$  alkyl; n is an integer from 1 to 4;  $R^8$  and  $R^9$  are each H or taken together form a double bond between C-22 and C-23; and pharmaceutically acceptable salts thereof, i.e., the bisphosphonate is linked at the C-25 position of the vitamin D moiety.

Also provided are conjugates of formula (X):

wherein  $R^1$  is H or OH;  $R^2$  is H or OH;  $R^3$  is  $CH_3$  or H;  $R^4$  s H or OH, X is O or S; Y is NH, O or NR wherein R is H or  $C_1$ - $C_4$  alkyl;  $R^5$  is H or OH; n is an

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integer from 1 to 4; R<sup>8</sup> and R<sup>9</sup> are each H or taken together form a double bond between C-22 and C-23; and pharmac utically acceptable salts thereof, i.e., the bisphosphonate is linked to the vitamin D moiety at C-3.

Also provided are conjugates of formula (XI):

$$R^{3}$$
 $R^{4}$ 
 $R^{5}$ 
 $R^{8}$ 
 $R^{8}$ 
 $R^{8}$ 
 $R^{5}$ 
 $R^{5}$ 
 $R^{6}$ 
 $R^{8}$ 
 $R^{8}$ 
 $R^{5}$ 
 $R^{6}$ 
 $R^{7}$ 
 $R^{7}$ 
 $R^{5}$ 
 $R^{5$ 

wherein  $R^2$  is H or OH;  $R^3$  is  $CH_3$  or H;  $R^4$  is H or OH, X is O or S; Y is NH, O or NR wherein R is H or  $C_1$ - $C_4$  alkyl;  $R^5$  is H or OH, n is an integer from 1 to 4,  $R^8$  and  $R^9$  are each H or taken together form a double bond between C-22 and C-23; and pharmaceutically acceptable salts thereof, i.e., the bisphosphonate linkage is at C-1 of the vitamin D moiety.

It is noted that typically the linkage between the bisphosphonate moiety and the vitamin D moiety is through a hydroxyl on the vitamin D where the hydroxyl is converted to a

group and is linked to the amine or hydroxy group, i.e., Y, of the bisphosphonate to form a carbamate-type or carbonate-type linkage. X can be O or S. For example, a hydroxyl group may be contained in the vitamin D structure at C-1, C-3, C-24, C-25, and conjugation can be effected at any hydroxyl position but is suitably one of the above.



### INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference		f Transmittal of International Search Report 20) as well as, where applicable, item 5 below.
2833.58	ACTION	
International application No.	International filing date (day/month/year)	(Earliest) Priority Date (day/month/year)
PCT/US 98/02899	13/02/1998	13/02/1997
Applicant		
BONE CARE INTERNATIONAL,	INC. et al.	· .
	een prepared by this International Searching Auth transmitted to the International Bureau.	nority and is transmitted to the applicant
, ,,		
This International Search Report consis		
X It is also accompanied by a co	opy of each prior art document cited in this report.	
		<del>and the second of the second </del>
1. Certain claims were found u	ınsearchable(see Box I).	
2. Unity of invention is lacking	ı(see Box II).	
•		
	contains disclosure of a nucleotide and/or amino	o acid sequence listing and the
	ed out on the basis of the sequence listing	
	ed with the international application. Irnished by the applicant separately from the inter	rnational application
	but not accompanied by a statement to the	
	matter going beyond the disclosure in the	
□ +	annesit and has the Australia	
<b>□</b> "	ranscribed by this Authority	
4. With regard to the title, X the	e text is approved as submitted by the applicant	
th	e text has been established by this Authority to re	ead as follows:
		·
5. With regard to the abstract,	o toyt is approved as submitted by the applicant	
	e text is approved as submitted by the applicant to text has been established, according to Rule 38	8.2(b) by this Authority as it appears in
во	ox III. The applicant may, within one month fromt earch Report, submit comments to this Authority.	the date of mailing of this International
	sarch rieport, submit comments to this Authority.	· ·
	•	
6. The figure of the <b>drawings</b> to be pu	•	[ Name of No. 12.
	s suggested by the applicant.	X None of the figures.
<b>=</b> .	ecause the applicant failed to suggest a figure. ecause this figure better characterizes the invention	on.
	readse and righte better characterizes the inventor	on.

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61K47/48 A61K31/59

According to International Patent Classification (IPC) or to both national classification and IPC

#### B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 \_\_A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

·	. DOCL	MENIS	COMPINE	HED H	O RE HEL	EVAN I

Category <sup>3</sup>	Citation of document, with indication, where appropriate, of the relevant passages	Retevant to claim No.
X,P	US 5 691 328 A (PETERSON ANDREW C ET AL) 25 November 1997 * cf. abstract, col. 6, lines 37-47, claims*	1-40
<b>X</b>	US 5 232 836 A (BOUILLON ROGER ET AL) 3 August 1993 *cf. abstract, col. 2, last para., col. 27, lines 25-35, col. 29, lines 20-27*	1-40
X	WO 93 07883 Å (ISIS PHARMACEUTICALS INC) 29 April 1993 *cf. claims 1-3*	1-40
·	-/	

X Further documents are listed in the continuation of box C.	χ Patent family members are listed in annex.
<ul> <li>Special categories of cited documents:</li> <li>"A" document defining the general state of the art which is not considered to be of particular relevance</li> <li>"E" earlier document but published on or after the international filing date</li> <li>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publicationdate of another citation or other special reason (as specified)</li> <li>"O" document referring to an oral disclosure, use, exhibition or other means</li> <li>"P" document published prior to the international filing date but later than the priority date claimed</li> </ul>	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention  "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone  "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.  "&" document member of the same patent family
Date of the actual completion of theinternational search	Date of mailing of the international search report
28 May 1998	16/06/1998
Name and mailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2  NL - 2280 HV Rijswijk  Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  Fax: (+31-70) 340-3016	Authorized officer  Stoltner, A

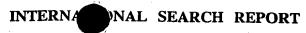
Form PCT/ISA/210 (second sheet) (July 1992)

### NTERNATI L SEARCH REPORT

PCT/US 98/02899

Category '	tion) DOCUMENTS CONSIDERED TO BE RELEVANT  Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
K	LONDOWSKI J.M., KOST S.B. ET AL.: "Biological activity of the C-1, C-3, C-25, betaD.glucopyranosides of 1,25-dihydroxyvitamin_D3"	1-40
	J. PHARMACOL. EXP. THER., vol. 237, no. 3, 1986, pages 837-840, XP002066239 * see the whole document*	

1



Information on patent family members

PCT/US 98/02899

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5691328 A	25-11-97	NONE	
US 5232836 A	03-08-93	FR 2631025 A EP 0341158 A ES 2045484 T JP 2262555 A US 5093519 A	10-11-89 08-11-89 16-01-94 25-10-90 03-03-92
WO 9307883 A	29-04-93	AU 2916292 A CA 2122030 A,C EP 0724447 A JP 6510791 T US 5578718 A	21-05-93 29-04-93 07-08-96 01-12-94 26-11-96